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**UTILITY
PATENT APPLICATION
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(Only for new nonprovisional applications under 37 CFR 1.53(b))

Attorney Docket No. 7547

First Inventor or Application Identifier

Raymond Edward Bolich, Jr.

Title

AEROSOL HAIR STYLING COMPOSITIONS CONTAINING SELECTED
POLYALKYLENE GLYCOL STYLING AGENTS

Express Mail Label No.

E J358988232US

APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents.

Assistant Commissioner for Patents
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1. ☒ Fee Transmittal Form
(Submit an original and a duplicate for fee processing.)
2. ☒ Specification [Total Pages = 27+ abstract]
(preferred arrangement set forth below)
 - Descriptive title of the Invention
 - Cross References to Related Applications
 - Statement Regarding Fed Sponsored R&D
 - Reference to Microfiche Appendix
 - Background of the Invention
 - Brief Summary of the Invention
 - Brief Description of the Drawings (if filed)
 - Detailed Description
 - Claim(s)
 - Abstract of the Disclosure
3. ☐ Drawing(s) (35 USC 113) [Total Sheets = 0]
4. ☐ Oath or Declaration [Total Pages = 2]
 - a. ☒ Newly executed (original or copy)
 - b. ☐ Copy from a prior application (37 CFR 1.63(d))
(for continuation/divisional with Box 17 completed)
[Note Box 5 below]
 - i. ☐ **DELETION OF INVENTOR(S)**
Signed statement attached deleting inventor(s)
named in the prior application, see 37 CFR
§§1.63(d)(2) and 1.33(b).
5. ☐ Incorporation By Reference (useable if Box 4b is checked)
The entire disclosure of the prior application, from
which a copy of the oath or declaration is supplied
under Box 4b, is considered as being part of the
disclosure of the accompanying application and is
hereby incorporated by reference therein.

6. ☐ Microfiche Computer Program (Appendix)
7. Nucleotide and/or Amino Acid Sequence Submission
(if applicable, all necessary)
 - a. ☐ Computer Readable Copy
 - b. ☐ Paper Copy (identical to computer copy)
 - c. ☐ Statement verifying identity of above copies

ACCOMPANYING APPLICATION PARTS

8. ☐ Assignment Papers (cover sheet & document(s))
9. ☐ 37 CFR 3.73(b) Statement ☒ Power of
(when there is an assignee) Attorney
10. ☐ English Translation Document (if applicable)
11. ☐ Information Disclosure ☐ Copies of
Statement (IDS)/PTO-1449 IDS Citations
12. ☐ Preliminary Amendment
13. ☒ Return Receipt Postcard (MPEP 503)
(Should be specifically itemized)
14. ☐ Small Entity ☐ Stmt. filed in prior application,
Statement(s) Status still proper and desired
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☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior application No. /

Prior application information: Examiner: _____ Group/Art Unit: _____

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(Insert Customer No. Or attach bar code label)

or ☒ Correspondence address below

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PTO/SB/29 (12/97)

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TOTAL CLAIMS (37 CFR 1.16(e))		40-20=	20	x \$18.00 =	\$360.00
INDEPENDENT CLAIMS (37 CFR 1.16(b))		4-3=	1	x \$78.00 =	\$78.00
MULTIPLE DEPENDENT CLAIMS (if applicable)(37 CFR 1.16(d))				+ \$260.00 =	\$0
				BASIC FEE (37 CFR 1.16(a))	\$760.00
				Total of above Calculations =	\$1198.00
Reduction by 50% for filing by small entity (Note 37 CFR 1.9, 1.27, 1.28).					\$0
				TOTAL =	\$1198.00

19. Small entity status:

- a. ☐ A small entity statement is enclosed.
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20. The Commissioner is hereby authorized to credit overpayments or charge the following fees to Deposit Account No. 16-2480:

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23. SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT REQUIRED

NAME	Joan B. Tucker (43,962)
SIGNATURE	<i>Joan B. Tucker</i>
DATE	May 5, 1999

[Page 2 of 2]

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43.962

Agent mailing application

Reg. No.

Signature of Agent mailing application

Joan B. Tucker

Aerosol Hair Styling Compositions Containing Select Polyalkylene Glycol Styling Agents

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Kenneth Wayne Rigney

Alic Anthony Scott

Case No. 7547

FIELD OF INVENTION

The present invention relates to aerosol hair styling compositions which provide improved hair restyling performance. In particular, the present invention relates to aerosol hair styling compositions which contain select polyalkylene glycols that provide for improved dry hair restyling performance without reapplication of the composition and/or any additional hair styling aids.

BACKGROUND OF THE INVENTION

Hair styling compositions are well known and are commercially available in a variety of forms including mousses, gels, lotions, pumps, or hairsprays. Many of these products contain various hair styling agents to provide temporary hair styling benefits such as body, hold, luster, improved hair feel, and good style retention.

Within this product group, hairsprays have become especially popular among consumers as an effective means for styling or "setting" the hair. The hairspray compositions typically comprise gums or resins styling agents to provide for good style retention and hair feel benefits. Most of the hairsprays, however, are sprayed onto the hair after the hair is styled in order to temporarily secure the desired hair style.

One method of providing temporary hair styling benefits from a styling product while styling the hair involves the use of a hair mousse. Many hair mousse products have been formulated such that the hair mousse composition can be applied to wet or damp hair before styling or "setting" the hair. Typically, the hair mousse compositions are aqueous formulations which contain water-soluble styling agents that provide adhesive properties to the hair while the hair is wet and being styled. These styling agents, however, can form hard breakable films on the hair as the styling process is near completion, and this can result in an unacceptable hair style or in a hair style that cannot be restyled unless additional water and/or supplemental styling products are added to the hair.

Moreover, the use of aqueous hair mousse compositions which contain water-soluble styling agents can leave the hair feeling unduly sticky and stiff.

One attempt at producing an aqueous hair mousse composition that is especially effective in providing good hair styling performance without leaving the hair feeling unduly stiff or sticky involves the formulation of an aqueous hair mousse which comprises a water-soluble styling polymer, a silicone emulsion, a liquid carrier comprising a combination of water and lower alcohols, and a silicone-linear polyoxyalkylene block copolymer surfactant which maintains stability of the silicone emulsion. Although these aqueous hair mousse compositions are effective in providing good hair styling performance and improved hair feel, they are also typically applied to wet or damp hair, and often lead to the application of an additional styling product once the hair has dried in order to achieve the desired final hairstyle form.

Another attempt at producing an aqueous aerosol hair mousse composition that provides good hair styling performance and improved hair feel involves the use of a combination of panthenol and certain low molecular weight polyalkylene glycols. The low molecular weight polyalkylene glycols, i.e., polyethylene glycols or polypropylene glycols having from 3 to about 12 ethylene glycol or propylene glycol units, help to effectively and efficiently deliver the panthenol to the hair. These hair mousse compositions, however, are also typically applied to wet or damp hair to achieve the desired hair conditioning benefits and generally require reapplication of the composition or another styling product to maintain or modify the original hairstyle.

Therefore, a need exists for hair styling compositions which provide for a desired hair style to be achieved and maintained, and which can be applied or sprayed on the hair during the styling process. A recent method of making a hair styling composition that can be applied to wet and/or dry hair during the styling process, and provide improved styling performance is described in JP 8-346608, published June 23, 1998. The hair styling compositions disclosed in this publication contain polyalkylene glyceryl ethers styling agents to provide for sustained hair styling performance and improved aesthetics such as longer moisture and luster to the hair. These polyalkylene glyceryl ether styling agents typically do not readily penetrate into the hair and can remain on the hair fibers to provide the improved hair styling benefits. The polyalkylene glyceryl ether containing compositions disclosed in this particular publication, however, tend to provide minimal or no hold to the hair, and this can result in poor style achievement and poor style retention performance.

Another recent method of making a hair styling composition that can be applied during the styling process, and provide improved styling performance is described in WO 98/38969, published September 11, 1998. The compositions disclosed in this publication use certain styling agents which deliver hair style performance to dry hair, and provide for the dry hair to be restyled without having to reapply the composition and without requiring the use of another styling product. The styling agents described in this reference include anionic, cationic, amphoteric, and nonionic styling polymers, preferably sulfonated anionic styling polymers which have an average molecular weight of

from about 500 to about 5,000,000. These styling agents, however, have exceptional cohesive strength which provides for the hair fibers to be firmly held together, and this can cause the dry hair to feel coarse and be difficult to comb, style, and restyle.

It has now been found that other hair styling compositions, particularly aerosol hair styling compositions, can be formulated to provide improved dry hair restyle performance without the need to reapply the composition and/or add any other styling aids. Unlike the compositions described in the WO 98/38969 publication, these new aerosol hair styling compositions comprise low molecular weight polyalkylene glycol styling agents that are liquids or semisolids under ambient conditions and that can remain as a liquid or semisolid after the compositions have been applied and allowed to dry on the hair. These select polyalkylene glycols can provide for a fluid film to be left on the hair which can be characterized as a reformable weld that allows the hair fibers to be separated by forces such as wind, and then re-adhere using styling techniques such as combing, brushing, or running your fingers through the hair. The ability of these select polyalkylene glycol styling agents to remain as a liquid or semisolid also provides for an aerosol hair styling product that can be applied and evenly distributed onto dry hair, resulting in the dry hair feeling extremely smooth and very easy to restyle. This improved restyle performance provided by these select polyalkylene glycols can last for extended periods of several days without the need to reapply the composition and/or add any additional styling aids on the hair.

It is therefore an object of the present invention to provide an aerosol hair styling composition which provides improved dry hair restyling performance and improved hair feel, and which contains select polyalkylene glycol styling agents that have a number average molecular weight of from about 190 to about 1500. It is also an object of the present invention to provide an aerosol hair styling composition in the form of a hair mousse which provides improved dry hair restyling performance and improved hair feel, and which contains select polyalkylene glycol styling agents that have a number average molecular weight of from about 190 to about 1500. It is yet another object of the present invention to provide an aerosol hair styling composition which provides improved dry hair restyling performance and improved hair feel for extended periods of time without the need to reapply the composition or add any other styling aids. It is still yet another object of the present invention to provide an aerosol hair styling composition that is a leave-on hair styling composition, and further to provide such a composition that does not feel unduly sticky or stiff after the composition has been applied and allowed to dry on the hair.

SUMMARY OF THE INVENTION

The present invention is directed to aerosol hair styling compositions which comprise (a) from about 5% to about 90% by weight of a polyalkylene glycol that is substantially free of polyalkylene glyceryl ethers and that has a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the

repeating alkylene oxide radicals has from 2 to 6 carbon atoms; (b) from about 1% to about 90% by weight of a liquid carrier; and (c) from about 5% to about 40% by weight of a propellant.

The present invention is also directed to hair mousse compositions which comprise (a) from about 5% to about 25% by weight of a polyalkylene glycol that is substantially free of polyalkylene glyceryl ethers and that has a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms; (b) from about 10% to about 90% by weight of water; and (c) from about 5% to about 40% by weight of a propellant. The present invention is also directed to a method of making such a hair mousse composition.

The present invention is also directed to hair styling compositions which comprise (a) from about 65% to about 99% by weight of a polyalkylene glycol that is substantially free of polyalkylene glyceryl ethers and that has a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms; and (b) from about 1% to about 30% by weight of a liquid carrier.

It has been found that the aerosol hair styling compositions of the present invention can provide improved dry hair restyling performance for several days without the need to reapply the composition or add any other styling aid on the hair. This is accomplished by formulating an aerosol hair styling composition, especially a hair mousse, that contains select polyalkylene glycol styling agents which are capable of being deposited on the hair as a liquid or semisolid and can remain a liquid or semisolid after the composition has been allowed to dry on the hair. These select styling agents can create a fluid film on the hair in the form of a reformable weld that provides for improved hair restyling performance such as 1) dry hair styled and restyled without reapplication of the compositions and/or water and/or any other styling aid, 2) dry hair restyled for several days by simply combing or brushing the hair, and 3) improved styling aesthetics including frizz and volume control, superior hair hold, and superior hair feel.

DETAILED DESCRIPTION OF THE INVENTION

The aerosol hair styling compositions of the present invention comprise select polyalkylene glycols that can be deposited on the hair and form reformable welds on the hair fibers. These compositions are intended for application to dry hair, and are preferably formulated as leave-on aerosol hair styling products.

The term "leave-on" as used herein refers to compositions that contain ingredients that are intended to be deposited and left on the hair for extended periods (e.g., several hours or days) until the ingredients are subsequently removed by water and/or by shampooing the hair.

The term "reformable weld" as used herein refers to residues which are left on dry hair and which contain materials that are liquid or semisolid at ambient condition, and that can remain as a

liquid or semisolid after the compositions described herein have been applied and allowed to dry on the hair.

The term "ambient conditions" as used herein refers to surrounding conditions at about one atmosphere of pressure, at about 50% relative humidity, and at about 25°C.

5 The term "substantially free" as used herein, unless otherwise specified, refers to preferred negative limitations of the compositions of the present invention, and are directed to the amount and concentration of polyalkylene glyceryl ether styling agents, or derivatives thereof, in the compositions. The term "substantially free" means that the compositions preferably contain less than an effective amount of such agents when used alone to provide any hair styling performance when
10 the compositions are applied to the hair. In this context, the negative limitations pertain only to those polyalkylene glyceryl ether styling agents which are also a liquid or semisolid under ambient conditions, and which are not silicone-containing materials. Generally, the compositions preferably contain less than 5%, more preferably less than 2%, even more preferably less than 1%, most preferably zero percent, of such agents by weight of the compositions.

15 The aerosol hair styling compositions of the present invention can comprise, consist of, or consist essentially of the essential elements and limitations of the invention described herein, as well as any of the additional or optional ingredients, components, or limitations described herein.

 All percentages, parts and ratios are by weight of the total composition, unless otherwise specified. All such weights as they pertain to listed ingredients are based on the active level and,
20 therefore, do not include solvents or by-products that may be included in commercially available materials, unless otherwise specified.

Styling Agent

25 The aerosol hair styling compositions of the present invention comprise a liquid or semisolid hair styling agent suitable for being left on dry hair as a liquid or semisolid after the composition has been applied and allowed to dry on the hair. These hair styling agents provide for a fluid film to be left on the hair which can be characterized as a reformable weld that provides dry hair restyling performance without the need to reapply the compositions or add additional styling aids on the hair.

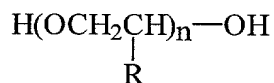
 The concentration of the styling agent may vary with each selected hair styling formulation,
30 but such concentrations will generally range from about 1% to about 90%, preferably from about 5% to about 75%, more preferably from about 7.5% to about 50%, even more preferably from about 10% to about 25%, by weight of the composition. For aerosol hair styling compositions in the form of a mousse product, the concentration of the styling agent preferably range from about 5% to about 25%, more preferably from about 7.5% to about 25%, even more preferably from about 10% to about 20%,
35 by weight of the composition.

 Suitable styling agents for use in the aerosol hair styling compositions of the present invention include any known or otherwise effective hair styling agents, other than polyalkylene

glyceryl ethers, that are liquids or semisolids under ambient conditions and that can remain a liquid or semisolid after the composition has been applied and allowed to dry on dry hair. It has been found that certain liquid or semisolid styling agents, particularly low molecular weight polyalkylene glycols, can leave a fluid film on the hair that allows the hair fibers to be separated by forces such as wind, and then re-adhere using styling techniques such as combing, brushing, or running your fingers through the hair. This separation/readherence property provided by the styling agents defined herein results in improved dry hair restyling performance for several days without leaving the hair feeling unduly sticky or stiff, and without having to reapply the compositions described herein and/or add any other additional styling aids on the hair.

Nonlimiting examples of styling agents suitable for use in the aerosol hair styling compositions of the present invention include water-soluble materials such as polyalkylene glycols, polyethylene/polypropylene glycol copolymers, polyethylene/polypropylene diol copolymers, polyglycerins, and mixtures thereof, and/or their derivatives, and/or mixtures thereof, excluding the water-soluble polyalkylene glyceryl ethers which are also liquids or semisolids under ambient conditions. In this context, the term "water-soluble" refers to those styling materials that have a solubility in water at 25°C of greater than 0.6%, preferably greater than 1.0%, more preferably greater than about 1.5% by weight.

Preferred styling agents suitable for use herein include those water-soluble polyalkylene glycols which conform to the formula:



wherein R is selected from the group consisting of H, methyl, and mixtures thereof. When R is H, these materials are polymers of ethylene oxide, which are also known as polyethylene oxides, polyoxyethylenes, and polyethylene glycols. When R is methyl, these materials are polymers of propylene oxide, which are also known as polypropylene oxides, polyoxypropylenes, and polypropylene glycols. When R is methyl, it is also understood that various positional isomers of the resulting polymers can exist.

In the above structure, n has an average value of from 4 to about 35, preferably from about 5 to about 35, more preferably from about 5 to about 30, and even more preferably from about 5 to about 20.

Specific examples of preferred polyalkylene glycol polymers include polyethylene/polypropylene glycol copolymers (e.g., methoxy, ethoxy, propoxy, butoxy, and pentoxy, polyethylene/polypropylene glycols), triglycerin, hexaglycerin, PPG-4, PPG-6, PEG-5, PEG-6, PEG-8, PEG-12, PEG-14, PEG-18, PEG-20, PEG-32, and mixtures thereof. Most preferred are those polyalkylene glycols which have a number average molecular weight of from about 190 to about 1500, preferably from about 300 to about 1200, more preferably from about 400 to about 1000; and from about 5 to about 35, preferably from about 5 to about 30, more preferably from about 5 to

about 20, repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms. Specific examples of the most preferred polyalkylene glycols include, but are not limited to, PPG-4 wherein R equals methyl and n has an average value of about 4; PEG-8 wherein R equals H and n has an average value of about 8 (PEG-8 is also known as Carbowax 400, which is available from Union Carbide); PEG-12 wherein R equals H and n has an average value of about 12 (PEG-12 is also known as Carbowax 600, which is available from Union Carbide); and PEG-20 wherein R equals H and n has an average value of about 20 (PEG-20 is also known as Carbowax 900, which is available from Union Carbide).

In addition to the styling agent, the aerosol hair styling compositions of the present invention may further comprise one or more optional styling polymers which can help provide improved initial hair hold performance. The total concentration of such optional styling polymers ranges from about 0.25% to about 5%, preferably from about 0.5% to about 2.5%, by weight of the compositions.

Optional styling polymers for use in combination with the styling agent defined herein include any known or otherwise effective styling polymer, provided that the optional styling polymer is soluble in the liquid carrier described herein which contains the optional styling polymer and styling agent, and provided that under test conditions of 27°C and 15% relative humidity the optional styling polymer is insoluble in the residue described herein and can form a solid film that is surrounded by the styling agent material after evaporation of the liquid carrier and any other volatile materials contained in the aerosol hair styling compositions of the present invention. Such optional styling polymers include, but are not limited to, polysaccharide styling polymers. Specific nonlimiting examples of suitable polysaccharide styling polymers include anionic polysaccharides, cationic polysaccharides, and glucosamine polysaccharide derivatives. The glucosamine polysaccharide derivatives are the preferred optional styling polymers.

Suitable optional cationic polysaccharide styling polymers for use herein include, but are not limited to, copolymers of hydroxyethylcellulose and diallyldimethyl ammonium chloride (referred to in the industry by CTFA as Polyquaternium-4) such as those commercially available from National Starch (Bridgewater, New Jersey) under the CELQUAT tradename (e.g., CELQUAT L-200 and CELQUAT H-100); and cationic quaternary ammonium-containing polymers, including, for example, homopolymers of hydroxyethyl cellulose reacted with a trimethyl ammonium substituted epoxide, (referred to in the industry by CTFA as Polyquaternium-10) such as those commercially available from Amerchol Corp. (Edison, New Jersey) under the UCARE tradename (e.g. UCARE POLYMER JR-400, and UCARE POLYMER LR-400), and those commercially available from National Starch (Bridgewater, New Jersey) under the CELQUAT tradename (e.g., CELQUAT SC 230 and CELQUAT SC 240).

Optional preferred polysaccharide styling polymers for use in combination with the styling agent in the aerosol hair styling compositions of the present invention include those polysaccharide

styling polymers which are derived from chitin, a glucosamine polysaccharide that is extracted from the shells of crabs, lobsters, and the like. An example of the use of chitin to make a chitin derivative for use in the compositions herein is the preparation of Chitosan, a water-soluble chitin derivative that is prepared by the known process of deacylation of the chitin compound. The chitin derivatives
 5 can also be prepared by other methods well known in the art for the preparation of such materials, including the hydroxypropylation of the chitin compound. The chitin derivatives suitable for use as an optional styling polymer herein include those chitin derivatives which are commercially available in a neutralized or unneutralized form. In the event that a neutralized chitin derivative is used, suitable neutralizing agents include, but are not limited to, lactic acid, pyrrolidone carboxylic acid,
 10 and glycolic acid.

Specific examples of preferred chitin derivatives for use as an optional styling polymer include, but are not limited to, Kytamer L and Kytamer PC (both are neutralized Chitosan materials commercially available from the Amerchol Corp., located in Edison, New Jersey); and Hydagen HCMF molecular weight (MW) of 50,000 to 1,000,000, Hydagen DCMF MW of 300,000 to
 15 2,000,000, and Hydagen CMFP MW of 500,000 to 5,000,000 (all are unneutralized Chitosan materials commercially available from the Henkel Corp., located in Hoboken, New Jersey). Kytamer L is the most preferred chitin derivative.

The hair styling compositions of the present invention can also be formulated as nonaerosol hair styling compositions which are suitable for topical application to the hair or skin. For
 20 nonaerosol hair styling product forms such as pomades, the concentration of the styling agent preferably ranges from about 65% to about 99%, more preferably from about 65% to about 95%, even more preferably from about 65% to about 90%, by weight of the composition. Pomades may further comprise combinations of the styling agent and optional styling polymers described herein, and any of the other optional ingredients described herein.

25

Liquid Carrier

The aerosol hair styling compositions of the present invention comprise any known or otherwise effective liquid carrier that is suitable for use in aerosol formulations intended for topical application to human hair or skin. The liquid carrier helps to solubilize or disperse the styling agents
 30 described hereinbefore. The liquid carrier can comprise one or more liquid carriers provided that the selected styling agent is sufficiently miscible or dispersible in the selected liquid carrier. The liquid carrier described herein is also suitable for use in the nonaerosol hair styling compositions of the present invention.

The total concentration of the liquid carrier in the composition will vary with the type of
 35 liquid carrier selected, the type of styling agent used in combination with the liquid carrier, and the solubility of the selected styling agent in the selected liquid carrier, and so forth. Preferred total

concentration of the liquid carrier ranges from about 1% to about 90%, preferably from about 3% to about 90%, more preferably from about 5% to about 85%, by weight of the composition.

Suitable liquid carriers for use in the aerosol hair styling compositions of the present invention include volatile liquid carrier materials. In this context, the term "volatile" refers to materials which have a boiling point of less than about 260°C, preferably from about 50°C to about 260°C, more preferably from about 60°C to about 200°C (at about one atmosphere of pressure).

Nonlimiting examples of volatile liquid carriers include water; organic solvents such as C₁-C₆ alkanols, carbitol, and acetone; and combinations thereof. Specific examples of suitable C₁-C₆ alkanols include, but are not limited to, ethanol, n-propanol, isopropanol, n-butanol, amyl alcohol, and mixtures thereof. Preferred C₁-C₆ alkanols include C₂-C₄ monohydric alcohols such as ethanol, isopropanol, and mixtures thereof.

In the event that the aerosol hair styling compositions of the present invention comprise combinations of water and an organic solvent such as C₁-C₆ alkanols, water is preferably included at concentrations of from about 40% to about 90%, more preferably from about 50% to about 90%, even more preferably from about 60% to about 90%; and the C₁-C₆ alkanols are preferably included at total concentrations of from about 1% to about 15%, more preferably from about 3% to about 15%, even more preferably from about 5% to about 10%, by weight of the composition.

Propellant

The aerosol hair styling compositions of the present invention comprise a propellant suitable for aerosol delivery of the composition to the desired application surface. The total concentration of the propellant in the aerosol hair styling composition can include one or more propellants, the total propellant concentration typically ranging from about 5% to about 40%, preferably from about 5% to about 25%, more preferably from about 5% to about 15%, by weight of the composition.

Nonlimiting examples of suitable propellants include hydrocarbons, nitrogen, carbon dioxide, nitrous oxide, atmospheric gas, 1,2-difluoroethane (Hydrofluorocarbon 152A) supplied as Dymel 152A by Dupont, dimethylether, and mixtures thereof. Preferred are the hydrocarbon propellants, specific examples of which include propane, butane, and isobutane. Most preferred is a hydrocarbon propellant containing a mixture of propane and isobutane, specific examples of which include Aeron A-46 and Aeron A-70 (both are commercially available from Diversified CPC).

Optional Components

In addition to the essential components described hereinbefore, the aerosol hair styling compositions of the present invention may further comprise one or more optional components known or otherwise effective for use in hair care or personal care products, provided that the optional components are physically and chemically compatible with the essential components described herein, or do not otherwise unduly impair product stability, aesthetics, or performance. Nonlimiting

examples of such optional components are disclosed in *International Cosmetic Ingredient Dictionary*, Fifth Edition, 1993, and *CTFA Cosmetic Ingredient Handbook*, Second Edition, 1992, both of which are incorporated by reference herein in their entirety. Some nonlimiting examples of such optional components are disclosed below.

5 Optional Gelling Agent

The aerosol hair styling compositions of the present invention preferably further comprise a gelling agent to help provide the desired viscosity to the residue which remains on the hair after the composition has been applied and allowed to dry on the hair. The preferred optional gelling agent also helps to provide for improved hair hold performance. Suitable optional gelling agents include
 10 any material known or otherwise effective in providing any gelling or measurable viscosity increase to the residue. The concentrations of the optional gelling agent in the compositions range from about 0.1% to about 10%, preferably from about 0.2% to about 5.0%, by weight of the compositions.

Nonlimiting examples of suitable optional gelling agents include crosslinked carboxylic acid polymers; unneutralized crosslinked carboxylic acid polymers; unneutralized modified
 15 crosslinked carboxylic acid polymers; crosslinked ethylene/maleic anhydride copolymers; unneutralized crosslinked ethylene/maleic anhydride copolymers (e.g., EMA 81 commercially available from Monsanto); unneutralized crosslinked allyl ether/acrylate copolymers (e.g., Salcare SC90 commercially available from Allied Colloids); unneutralized crosslinked copolymers of sodium polyacrylate, mineral oil, and PPG-1 trideceth-6 (e.g., Salcare SC91 commercially available from Allied Colloids); unneutralized crosslinked copolymers of methyl vinyl ether and maleic
 20 anhydride (e.g., Stabileze QM-PVM/MA copolymer commercially available from International Specialty Products); hydrophobically modified nonionic cellulose polymers; hydrophobically modified ethoxylate urethane polymers (e.g., Ucare Polyphobe Series of alkali swellable polymers commercially available from Union Carbide); and combinations thereof. In this context, the term
 25 "unneutralized" means that the optional polymer and copolymer gelling agent materials contain unneutralized acid monomers.

Preferred optional gelling agents include water-soluble unneutralized crosslinked ethylene/maleic anhydride copolymers, water-soluble unneutralized crosslinked carboxylic acid polymers, and water-soluble hydrophobically modified nonionic cellulose polymers. The
 30 crosslinked carboxylic acid polymers and hydrophobically modified nonionic cellulose polymers are described in detail hereinbelow.

Carboxylic Acid Polymers

The optional carboxylic acid polymers suitable for use herein are those crosslinked compounds containing one or more monomers derived from acrylic acid, substituted acrylic acids,
 35 esters of acrylic acid, esters of substituted acrylic acids, corresponding salts thereof, and combinations thereof, wherein the crosslinking agent contains two or more carbon-carbon double bonds and is derived from a polyhydric alcohol. Specific examples of these carboxylic acid

polymers include crosslinked carboxylic acid homopolymers and crosslinked carboxylic acid copolymers. Combinations of these two types of polymers are also useful herein.

The term "substituted" as used herein refers to chemical moieties known or otherwise effective for attachment to gelling agents or other compounds. Such substituents include those listed and described in C. Hansch and A. Leo, *Substituent Constants for Correlation Analysis in Chemistry and Biology* (1979), which listing and description are incorporated herein by reference. Examples of such substituents include, but are not limited to, alkyl, alkenyl, alkoxy, hydroxy, oxo, nitro, amino, aminoalkyl (e.g., aminomethyl, etc.), cyano, halo, carboxy, alkoxyaceyl (e.g., carboethoxy, etc.), thiol, aryl, cycloalkyl, heteroaryl, heterocycloalkyl (e.g., piperidinyl, morpholinyl, pyrrolidinyl, etc.), imino, thioxo, hydroxyalkyl, aryloxy, arylalkyl, and combinations thereof.

The term "corresponding salts" as used herein refers to cationic salts formed at any acidic (e.g., carboxyl) group, or anionic salts formed at any basic (e.g., amino) group, either of which are suitable for topical application to human skin. Many such salts are known in the art, examples of which are described in World Patent Publication 87/05297, Johnston et al., published September 11, 1987, which description is incorporated herein by reference.

Preferred optional crosslinked carboxylic acid polymers are those crosslinked carboxylic acid homopolymers or copolymers which contain unneutralized acid monomers. It has been found that crosslinked carboxylic acid polymers which have unneutralized acid monomers are especially effective in providing gelling properties to the residue without suppressing the ease of removability of the residue by shampooing the hair.

Partially or fully neutralized crosslinked carboxylic acid polymers are also suitable for use as an optional gelling agent in the hair styling compositions of the present invention, provided that these carboxylic acid polymers are included in combination with one or more styling agents which have an average solubility parameter of above about 14 (cal/cm³)^{0.5} to about 20 (cal/cm³)^{0.5}. Solubility parameters for the styling agents or other materials, and means for determining such parameters, are well known in the chemical arts. A description of solubility parameters and means for determining them are described by C.D. Vaughan, "Solubility Effects in Product, Package, Penetration and Preservation" 103 *Cosmetics and Toiletries* 47-69, October 1988; and C. D. Vaughan, "Using Solubility Parameters in Cosmetics Formulation", 36 *J. Soc. Cosmetic Chemists* 319-333, September/October, 1988, which descriptions are incorporated herein by reference.

Suitable crosslinked carboxylic acid homopolymers include those crosslinked homopolymers which have an acrylic acid monomer, or derivative thereof, (e.g., wherein the acrylic acid has substituents on the two and three carbon positions independently selected from the group consisting of C₁₋₄ alkyl, -CN, -COOH, and mixtures thereof). Preferred monomers include acrylic acid, methacrylic acid, ethacrylic acid, and mixtures thereof, with acrylic acid being the most preferred.

Suitable crosslinked carboxylic acid copolymers include those crosslinked copolymers which have a first monomer selected from the group consisting of an acrylic acid or derivative thereof as defined above, a short chain alcohol (i.e., a C₁₋₄) acrylate ester monomer or derivative thereof (e.g., wherein the acrylic acid portion of the ester has substituents on the two and three carbon positions independently selected from the group consisting of C₁₋₄ alkyl, -CN, -COOH, and mixtures thereof), and mixtures thereof; and a second monomer which is a long chain alcohol (i.e. C₈₋₄₀) acrylate ester monomer or derivative thereof (e.g., wherein the acrylic acid portion of the ester has substituents on the two and three carbon positions independently selected from the group consisting of C₁₋₄ alkyl, -CN, -COOH, and mixtures thereof). Preferred first monomers include acrylic acid, methacrylic acid, and ethacrylic acid, C₁₋₄ alcohol acrylate esters, C₁₋₄ alcohol methacrylate esters, C₁₋₄ alcohol ethacrylate esters, and mixtures thereof, with acrylic acid, methacrylic acid, C₁₋₄ alcohol acrylate esters, C₁₋₄ alcohol methacrylate esters, and mixtures thereof being the most preferred. Preferred second monomers include C₈₋₄₀ alkyl acrylate esters, with C₁₀₋₃₀ alkyl acrylate esters being the most preferred. In other words, the preferred crosslinked carboxylic acid copolymers include those copolymers which have a first monomer selected from the group consisting of acrylic acid, methacrylic acid, C₁₋₄ alcohol acrylate esters, C₁₋₄ alcohol methacrylate esters, and mixtures thereof, and a C₁₀₋₃₀ alkyl acrylate ester second monomer.

The crosslinking agent in both of these types of polymers is a polyalkenyl polyether of a polyhydric alcohol containing more than one alkenyl ether group per molecule, wherein the parent polyhydric alcohol contains at least 3 carbon atoms and at least 3 hydroxyl groups. Preferred crosslinking agents are those selected from the group consisting of allyl ethers of sucrose, allyl ethers of pentaerythritol, and mixtures thereof.

Examples of commercially available crosslinked carboxylic acid homopolymers suitable for use herein include the carbomers, which are homopolymers of acrylic acid crosslinked with allyl ethers of sucrose or pentaerythritol. The carbomers are available as the Carbopol® 900 series from B.F. Goodrich. Most preferred are the commercially available carbomers which have unneutralized acid monomers.

Examples of commercially available crosslinked carboxylic acid copolymers suitable for use herein include copolymers of C₁₀₋₃₀ alkyl acrylates with one or more monomers of acrylic acid, methacrylic acid, or one of their short chain (i.e. C₁₋₄ alcohol) esters, wherein the crosslinking agent is an allyl ether of sucrose or pentaerythritol. These copolymers are known as acrylates/C₁₀₋₃₀ alkyl acrylate crosspolymers and are commercially available as Carbopol® 1342, Pemulen TR-1, and Pemulen TR-2, from B.F. Goodrich.

Suitable crosslinked carboxylic acid polymers are more fully described in U.S. Patent No. 5,087,445, to Haffey et al., issued February 11, 1992; U.S. Patent No. 4,509,949, to Huang et al., issued April 5, 1985; U.S. Patent No. 2,798,053, to Brown, issued July 2, 1957; which descriptions

are incorporated by reference herein. See also, CTFA International Cosmetic Ingredient Dictionary, fourth edition, 1991, pp. 12 and 80; which description is also incorporated herein by reference.

Nonionic Cellulose Polymers

The preferred optional gelling agent suitable for use in the aerosol hair styling compositions of the present invention also include water-soluble hydrophobically modified nonionic cellulose polymers. The term "water-soluble hydrophobically modified nonionic polymers" refers to those water-soluble nonionic polymers which have been modified to comprise substituted hydrophobic groups to make the polymer less soluble in water. Hence, the nonionic cellulose polymers comprise a water-soluble cellulosic chain (or hydrophilic cellulosic chain) which forms the backbone, wherein the backbone comprise substituted hydrophobic groups. Suitable substituted hydrophobic groups include C8-C22 alkyl, arylalkyl, alkylaryl groups, and mixtures thereof. The degree of substitution on the backbone should be from about 0.10% to about 1.0%, depending on the particular polymer backbone. The nonionic cellulose polymers generally contain a ratio of hydrophilic substituents to hydrophobic substituents of from about 10:1 to about 1000:1.

Nonlimiting examples of preferred hydrophobically modified nonionic cellulose polymers include those nonionic cellulose polymers which comprise a cellulose ether substrate and a long chain alkyl modifier. In this context, the term "long chain alkyl modifier" means that the modifying compound can comprise an alkyl radical or other functional groups such as an aliphatic hydroxyalkyl radical, a urethane radical, or an acyl radical. These polymers, and methods of making the polymers, are also described in U.S. Patent 4,228,277, issued to Landoll on October 14, 1980, which description is incorporated herein by reference.

Suitable cellulose ether substrates include any known or otherwise effective water-soluble nonionic cellulose ether. Nonlimiting examples of suitable water-soluble nonionic cellulose ethers include, but are not limited to, hydroxyethyl cellulose, hydroxypropyl cellulose, methyl cellulose, hydroxypropyl methyl cellulose, ethyl hydroxyethyl cellulose, and methyl hydroxyethyl cellulose. The amount of the nonionic substituent (e.g., methyl, hydroxyethyl, or hydroxypropyl substituent) is not critical provided that the amount is sufficient to assure that the ether is water-soluble.

The preferred cellulose ether substrate is hydroxyethyl cellulose (HEC) which has a weight average molecular weight of from about 50,000 to about 700,000. It has been found that hydroxyethyl cellulose is the most hydrophilic suitable cellulose ether substrate, and therefore, can allow for greater modification than other suitable water-soluble cellulose ether substrates before water-insolubility is achieved. Accordingly, control of the modification process and control of the properties of the modified substrate can be more precise with hydroxyethyl cellulose substrates. Hydrophilicity of the most commonly used nonionic cellulose ethers ranges from hydroxyethyl to hydroxypropyl to hydroxypropylmethyl to methyl, with hydroxyethyl being the most hydrophilic and methyl being the least hydrophilic.

The long chain alkyl modifier can be attached to the cellulose ether substrate via an ether, ester, or urethane linkage. The ether linkage is preferred. The size and effect of the hydrocarbon chain of suitable modifiers effectively conceals any noticeable effect derived from functional groups other than alkyl halides which links the modifier to the substrate. In other words, if the long chain alkyl modifier is an epoxide containing an α -hydroxyalkyl radical, an isocyanate containing a urethane radical, or an acyl chloride containing an acyl radical, the effect of these functional groups are unnoticeable and the performance of the hydrophobically modified nonionic cellulose polymer is not significantly different from a polymer modified with an alkyl halide modifier.

Specific examples of preferred hydrophobically modified nonionic cellulose polymers include Natrosol Plus Grade 330, and Natrosol Plus CS Grade D-67, both commercially available from the Aqualon Company located in Wilmington, Delaware. Natrosol Plus Grade 330 is a hydrophobically modified hydroxyethylcellulose which has a weight average molecular weight of approximately 300,000 prior to modification; which has been substituted with from about 0.4% to about 0.8%, by weight, of an alkyl modifier having sixteen carbon atoms; and which has a hydroxyethyl molar substitution of from about 3.0 to about 3.7. Natrosol Plus CS Grade D-67 is a hydrophobically modified hydroxyethylcellulose which has a weight average molecular weight of approximately 700,000 prior to modification; which has been substituted with from about 0.50% to about 0.95%, by weight, of an alkyl modifier having sixteen carbon atoms; and which has a hydroxyethyl molar substitution of from about 2.3 to about 3.3.

Other cellulose polymers which can provide measurable viscosity increase to the residue are also suitable for use as an optional gelling agent herein. A specific example of other suitable cellulose polymers include a water-soluble hydrophobically modified cationic cellulose polymer commercially available as Quatrosol from Amerchol.

Optional Feel Enhancers

The aerosol hair styling compositions of the present invention may further comprise a feel enhancer to help provide improved dry hair feel. Concentrations of the optional feel enhancer will typically range from about 0.1% to about 15%, preferably from about 0.2% to about 10%, more preferably from about 0.5% to about 10%, by weight of the compositions.

Suitable optional feel enhancers include nonvolatile silicones, cationic polymers excluding the cationic polymers suitable for use as optional styling polymers or optional gelling agents herein, cationic surfactants, and mixtures thereof. These feel enhancers help to provide improved dry hair feel benefits such as nonsticky, unstiff, flexible, and natural feeling hair.

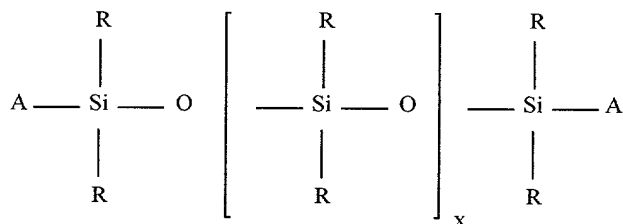
Nonlimiting examples of nonvolatile silicones suitable for use as an optional feel enhancer herein include nonvolatile soluble silicones, nonvolatile insoluble silicones, or combinations thereof. In this context, the term "soluble" means that the silicone feel enhancer is miscible with the liquid carrier so as to form part of the same phase. Conversely, the term "insoluble" means that the silicone feel enhancer forms a separate, discontinuous phase from the liquid carrier, such as in the form of an

emulsion, microemulsion, or a suspension of droplets of the silicone. The term "nonvolatile" as used in this context means that the silicone feel enhancer has a boiling point of at least about 265°C, preferably at least about 275°C, more preferably at least about 300°C. Such materials exhibit very low or no significant vapor pressure at ambient conditions.

5 Suitable nonvolatile soluble silicones include silicone copolyols, such as dimethicone copolyols, e.g. polyether siloxane-modified polymers, such as polypropylene oxide, polyethylene oxide modified polydimethylsiloxane, wherein the level of ethylene and/or propylene oxide is sufficient to allow solubility in the composition.

10 Suitable nonvolatile insoluble silicones include those insoluble silicone fluids such as polyalkylsiloxanes, polyarylsiloxanes, polyalkylaryl siloxanes, polyether siloxane copolymers, and mixtures thereof. Other nonvolatile insoluble silicone fluids having feel enhancing properties can also be used. In this context, the term "silicone fluid" refers to those silicone materials which have a viscosity of less than about 1,000,000 centistokes (cs) at 25°C. Generally, the viscosity of the fluid will be between about 5 cs and about 1,000,000 cs, preferably between about 10 cs and about 15 300,000 cs at 25°C. The viscosity can be measured by means of a glass capillary viscometer as set forth in Dow Corning Corporate Test Method CTM0004, July 20, 1970.

Nonlimiting examples of silicone fluids for use in the aerosol hair styling compositions of the present invention include polyalkyl or polyaryl siloxanes which conform to the formula:



20 wherein R is alkyl or aryl, and x is an integer from about 7 to about 8,000. "A" represents groups which block the ends of the silicone chains.

25 The alkyl or aryl groups substituted on the siloxane chain (R) or at the ends of the siloxane chains (A) may have any structure as long as the resulting silicones remain fluid at ambient conditions, are hydrophobic, are neither irritating, toxic nor otherwise harmful when applied to the hair or skin, are compatible with the other components of the composition, are chemically stable under normal use and storage conditions, and are capable of being deposited on and of improving the feel of hair or skin.

30 Suitable A groups include methyl, ethyl, phenyl, phenylene, hydroxy, methoxy, ethoxy, propoxy, and aryloxy. The two R groups on the silicone atom may represent the same group or different groups. Preferably, the two R groups represent the same group. Suitable R groups include methyl, ethyl, propyl, phenyl, methylphenyl and phenylmethyl. The preferred silicone fluids are polydimethylsiloxane, polydiethylsiloxane, and polymethylphenylsiloxane. Polydimethylsiloxane is most preferred.

Specific examples of nonvolatile insoluble silicone fluids include, but are not limited to, polydimethylsiloxanes such as the Viscasil R® and SF96® series (commercially available from the General Electric Co.), and the Dow Corning 200® series (commercially available from Dow Corning Corp.); polymethylphenylsiloxane such as SF1075® methyl phenyl fluid (commercially available from the General Electric Co.), and 556 Cosmetic Grade Fluid® (commercially available from Dow Corning Corp.); and a polypropylene oxide modified polydimethylsiloxane such as Dow Corning DC-1248®.

Other suitable silicone fluids are disclosed in U.S. Patent 2,826,551, issued to Geen; U.S. Patent 3,964,500, issued to Drakoff, on June 22, 1976; U.S. Patent 4,364,837, issued to Pader; and British Patent 849,433, issued to Woolston, all of which disclosures are incorporated herein by reference.

Other suitable silicone feel enhancers include insoluble silicone gum. The term "silicone gum", as used herein, means polyorganosiloxane materials having a viscosity at 25°C of greater than or equal to 1,000,000 centistokes. Silicone gums are described by Petrarch and others including U.S. Patent 4,152,416, Spitzer et al., issued May 1, 1979; and Noll, Walter, Chemistry and Technology of Silicones, New York: Academic Press 1968. The "silicone gums" will typically have a mass molecular weight in excess of about 200,000, generally between about 200,000 and about 1,000,000. Specific examples include polydimethylsiloxane, (polydimethylsiloxane) (methylvinylsiloxane) copolymer, poly(dimethylsiloxane) (diphenyl siloxane)(methylvinylsiloxane) copolymer, and mixtures thereof.

Other suitable silicone feel enhancers include a silicone resin. Silicone resins are highly crosslinked polymeric siloxane systems. The crosslinking is introduced through the incorporation of trifunctional and tetrafunctional silanes with monofunctional or difunctional, or both, silanes during manufacture of the silicone resin. As is well understood in the art, the degree of crosslinking that is required in order to result in a silicone resin will vary according to the specific silane units incorporated into the silicone resin. In general, silicone materials which have a sufficient level of trifunctional and tetrafunctional siloxane monomer units (and hence, a sufficient level of crosslinking) such that they dry down to a rigid, or hard, film are considered to be silicone resins. The ratio of oxygen atoms to silicon atoms is indicative of the level of crosslinking in a particular silicone material. Silicone materials which have at least about 1.1 oxygen atoms per silicon atom will generally be silicone resins herein. Preferably, the ratio of oxygen:silicon atoms is at least about 1.2:1.0. Silanes used in the manufacture of silicone resins include monomethyl-, dimethyl-, trimethyl-, monophenyl-, diphenyl-, methylphenyl-, monovinyl-, and methylvinyl-chlorosilanes, and tetrachlorosilane, with the methyl-substituted silanes being most commonly utilized. Preferred resins are offered by General Electric as GE SS4230® and SS4267®. Commercially available silicone resins will generally be supplied in a dissolved form in a low viscosity volatile or nonvolatile silicone fluid. The silicone resins for

use herein should be supplied and incorporated into the present compositions in such dissolved form, as will be readily apparent to those skilled in the art.

Background material on silicones including sections discussing silicone fluids, gums, and resins, as well as manufacture of silicones, can be found in Encyclopedia of Polymer Science and Engineering, Volume 15, Second Edition, pp 204-308, John Wiley & Sons, Inc., 1989, incorporated herein by reference.

Other Optional Materials

Other optional materials suitable for use in the aerosol hair styling compositions of the present invention include, but are not limited to, preservatives, surfactants, conditioning polymers, electrolytes, fatty alcohols, hair dyes, antidandruff actives, odor masking agents, pH adjusting agents, perfume oils, perfume solubilizing agents, sequestering agents, emollients, lubricants and penetrants such as various lanolin compounds, protein hydrolysates and other protein derivatives, sunscreens, volatile silicone fluids, and isoparaffins. The concentration of such optional ingredients generally ranges from zero to about 25%, more typically from about 0.05% to about 25%, even more typically from about 0.1% to about 15%, by weight of the composition.

Method of Manufacture

The aerosol hair styling compositions of the present invention may be prepared by any known or otherwise effective technique, suitable for providing an aerosol hair styling composition provided that the composition is formulated to contain the requisite hair styling agent defined herein.

Methods for preparing the aerosol hair styling compositions of the present invention include conventional formulation and mixing techniques. Suitable methods include combining the styling agent with the liquid carrier, and thoroughly mixing until the styling agent is homogeneously dispersed and dissolved in the liquid carrier. Any remaining ingredients such as perfume, the optional styling polymer, and the optional preferred gelling agent, can then be added and dispersed into the mixture. The resultant liquid mixture is then packaged into a suitable container such as an aerosol dispenser, and the propellant is then added.

The aerosol hair styling composition of the present invention can be contained or dispensed in any known or otherwise effective aerosol container or delivery system. All such containers or delivery systems should be compatible with the essential and any selected optional ingredients of the hair styling composition of the present invention.

Alternatively, pressurized aerosol dispensers can be used where the propellant is separated from contact with the hair styling composition by use of specialized containers such as a two compartment can of the type sold under the tradename SEPRO from American National Can Corp.

The hair styling compositions of present invention can also be formulated as nonaerosol compositions. Methods for preparing nonaerosol hair styling compositions such as pomades include

any known or otherwise effective manufacturing or formulation method for formulating such products.

Method Of Use

5 The hair styling compositions of the present invention are used to provide hairstyle/hold benefits without having to reapply the compositions for several days. An effective amount of the composition is either sprayed or applied onto dry hair before or after the hair is styled. As used herein "effective amount" means an amount sufficient to provide the hair hold and style performance desired according to the length and texture of the hair.

10

EXAMPLES

 The following examples further describe and demonstrate embodiments within the scope of the present invention. The examples are given solely for the purpose of illustration and are not to be construed as limitations of the present invention, as many variations thereof are possible without departing from the spirit and scope of the invention. All exemplified concentrations are weight-weight percents, unless otherwise specified.

15

Examples I-VIII

 The following Examples I-VIII describe hair mousse compositions of the present invention. Each of the exemplified compositions are prepared by combining all of the listed components, except the propellant, and mixing the combination for 40 minutes. The resultant liquid mixture is then filled into an aerosol container, and the propellant is added. Each of the exemplified hair mousse compositions provides improved dry hair restyle performance without the need to reapply the composition or add any other additional styling aids.

20

Hair Mousse Compositions

Component:	Example I	Example II	Example III	Example IV
PEG-8 ¹	20.0			
PEG-12 ²	---	15.0	---	---
PEG-20 ³	---		20.0	20.0
Ethanol (Denatured)	10.0	5.0	5.0	5.0
Polydimethylsiloxane (350 cs) ⁴	---	---	5.0	5.0
Silicone-Polyether Copolymer ⁵	5.0	---	---	---
Sodium Cocoyl Isethionate	---	0.50	---	---
Benzyl Alcohol	0.50	0.50	0.50	---
Undeceth-9 ⁶	---	0.50	0.50	---
Cocamide DEA	0.50	---	---	0.50
Lauramide DEA	---	---	---	0.50
Cocamidopropyl Betaine	0.50	---	---	---
DMDM Hydantoin	---	---	---	0.50
Phenoxyethanol	0.30	0.30	0.30	0.30
Perfume	0.15	0.15	0.15	0.15
Disodium EDTA	---	0.12	0.12	0.12
Propellant Aeron A-70 ⁷	7.0	10.0	---	---
Propellant Aeron A-46 ⁷	---	---	7.0	10.0
Distilled Water	qs	qs	qs	qs

1 - Carbowax 400 available from Union Carbide

2 - Carbowax 600 available from Union Carbide

3 - Carbowax 900 available from Union Carbide

4 - SF96-350 available from General Electric

5 - Silwet L-77 available from Witco

6 - Neodol 1-9 available from Shell

7 - Propane/Isobutane mixture available from Diversified CPC

Hair Mousse Compositions

Component:	Example V	Example VI	Example VII	Example VIII
PEG-6 ⁸	10.0	---	---	---
PEG-8 ¹	---	5.0	---	25.0
PEG-18	---	---	5.0	---
PEG-32 ⁹	10.0	---	---	---
Ethanol (Denatured)	5.0	---	---	5.0
DC 2-1559 Emulsion ¹⁰	---	10.0	0.15	---
Triglycerin	---	---	5.0	---
Glycerin	---	5.0	---	---
Dilauryl Acetyl Dimonium Chloride ¹¹	---	---	---	5.0
Ditalowdimonium Chloride	0.50	---	---	---
Benzyl Alcohol	---	---	---	0.50
Undeceth-9 ⁵	0.50	---	---	---
Phenoxyethanol	---	---	---	0.30
Steartrimonium Chloride	---	0.30	0.30	---
PEG-60 Castor Oil	---	0.20	0.20	---
Perfume	0.15	---	---	0.15
Disodium EDTA	---	---	---	0.09
Propellant Aeron A-70 ⁷	10.0	---	---	---
Propellant Aeron A-46 ⁷	---	10.0	10.0	7.0
Distilled Water	qs	qs	qs	qs

8 - Carbowax 300 available from Union Carbide

9 - Carbowax 1450 available from Union Carbide

10 - 60% silicone emulsion available from Dow Corning

11 - Schercoquat ALA available from Scher

5

Examples IX-XVII

- 10 The following Examples IX-XVII describe hair mousse compositions of the present invention. Each of the exemplified compositions are prepared by adding the styling agent to water and mixing for one hour until the styling agent is dissolved. The remaining ingredients, except the propellant, are then added and the resulting solution is stirred. The composition is then packaged into an aerosol container, and the propellant is added. Each of the exemplified hair mousse compositions provides improved dry hair restyle performance without the need to reapply the
- 15 composition or add any other additional styling aids.

Hair Mousse Compositions

Component:	Example IX	Example X	Example XI	Example XII
PEG-8 ¹	18.0	18.0	15.0	10.0
PEG-12 ²	---	---	---	---
PEG-20 ³	---	---	---	---
Kytamer L ¹²	---	1.80	---	---
EMA 81 ¹³	0.72	---	---	---
Laponite XLG ¹⁴	---	---	1.00	---
Hexaglycerol ¹⁵	---	---	---	10.0
Carbopol 934 ¹⁶	---	---	---	0.10
Permulan TR-1 ¹⁷	---	---	---	---
Benzyl Alcohol	0.45	0.45	0.50	0.50
Phenoxyethanol	0.27	0.27	0.30	0.30
Undeceth-9 ⁵	0.27	0.27	0.30	0.30
Sodium Cocoyl Isethionate	0.18	---	---	0.20
Disodium EDTA	0.11	---	0.12	0.12
Perfume	0.13	0.11	0.15	0.15
Aminomethylpropanol	---	---	---	0.10
Kathon CG ¹⁸	---	---	---	---
Propellant Aeron A-46 ⁷	10.0	---	7.50	10.0
Propellant Aeron A-70 ⁷	---	10.0	---	---
Distilled Water	qs	qs	qs	qs

12 - neutralized Chitosan available from Amerchol

13 - unneutralized ethylene/maleic anhydride copolymer available from Monsanto

14 - styling clay available from Southern Clay Products

15 - available from Solvay Interlox, Inc.

16 - crosslinked carboxylic acid homopolymer available from B. F. Goodrich

17 - crosslinked carboxylic acid copolymer available from B. F. Goodrich

18 - preservative available from Rohm & Haas

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Hair Mousse Compositions

Component:	Example XIII	Example XIV	Example XV	Example XVI	Example XVII
PEG-8 ¹	20.0	---	---	16.0	20.0
PEG-12 ²	---	12.50	---	---	---
PEG-20 ³	---	---	15.0	---	---
Sodium Carrageenan ¹⁹	---	---	---	1.50	---
Celquat SC 240 ²⁰	---	---	---	---	1.00
Carbopol 934 ¹⁶	0.50	0.50	---	---	---
Permulan TR-1 ¹⁷	---	---	0.50	---	---
Benzyl Alcohol	0.50	0.50	---	---	---
Phenoxyethanol	0.30	0.30	---	---	---
Undeceth-9 ⁵	0.30	0.15	0.30	---	---
Sodium Cocoyl Isethionate	---	0.15	0.20	0.40	0.35
Disodium EDTA	0.12	0.12	---	---	---
Perfume	0.15	0.15	0.15	0.12	0.25
Aminomethylpropanol	---	---	---	---	---
Kathon CG ¹⁸	---	---	0.04	0.04	0.04
Propellant Aeron A-46 ⁷	---	---	10.0	---	---
Propellant Aeron A-70 ⁷	10.0	10.0	---	---	---
Distilled Water	qs	qs	qs	qs	qs

19 - anionic polysaccharide polymer available from Aqualon

20 - cationic polysaccharide copolymer available from National Starch

Examples XVIII

The hair styling compositions of the present invention include the pomade embodiment described below. The pomade is formulated by dispersing the Carbopol 934 into a solution containing all the listed ingredients except the aminomethylpropanol, and thoroughly mixing for 30 minutes. The aminomethylpropanol is then slowly added to and thoroughly mixed in the resulting solution for 10 minutes. The resultant composition may be packaged into any well known container for packaging such pomade compositions. The exemplified pomade composition of the present invention provides improved dry hair restyle performance without the need to reapply the composition or add any other additional styling aids.

Pomade Composition

Component	Weight %
PEG-14	80.00
Carbopol 934 ¹⁶	1.00
Aminomethylpropanol	0.30
Polysorbate 80	0.40
Fragrance	0.20
Distilled Water	Q.S.

Examples XIX

The hair styling compositions of the present invention include the spray-on-gel embodiment described below. The spray-on-gel is formulated by combining all the listed ingredients, except the propellant, and mixing the combination for 10 minutes. The composition is then packaged into an aerosol container, and the propellant is added. The exemplified spray-on-gel composition of the present invention provides improved dry hair restyle performance without the need to reapply the composition or add any other additional styling aids.

Spray-on-gel Composition

Component	Weight %
PEG-20 ³	8.00
Fragrance	0.10
Isobutane	5.00
Dimethyl Ether	25.00
Ethanol, Denatured	20.00
Distilled Water	Q.S.

WHAT IS CLAIMED IS:

1. An aerosol hair styling composition comprising:
 - (a) from about 5% to about 90% by weight of a polyalkylene glycol that is substantially free of polyalkylene glyceryl ethers and that has a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms;
 - (b) from about 1% to about 90% by weight of a liquid carrier; and
 - (c) from about 5% to about 40% by weight of a propellant.
2. The composition of Claim 1 wherein the composition comprises from about 7.5% to about 50% by weight of the polyalkylene glycol.
3. The composition of Claim 2 wherein the polyalkylene glycol has a number average molecular weight of from about 400 to about 1500.
4. The composition of Claim 3 wherein the polyalkylene glycol is selected from the group consisting of ethoxy polyethylene/polypropylene glycol copolymers, methoxy polyethylene/polypropylene glycol copolymers, propoxy polyethylene/polypropylene glycol copolymers, butoxy polyethylene/polypropylene glycol copolymers, pentoxy polyethylene/polypropylene glycol copolymers, triglycerin, hexaglycerin, PPG-4, PPG-6, PEG-5, PEG-6, PEG-8, PEG-12, PEG-14, PEG-18, PEG-20, PEG-32, and mixtures thereof.
5. The composition of Claim 4 wherein the polyalkylene glycol is selected from the group consisting of PPG-4, PEG-8, PEG-12, PEG-20, and mixtures thereof.
6. The composition of Claim 2 wherein the liquid carrier is selected from the group consisting of water, C₁-C₆ alkanols, carbitol, acetone, and mixtures thereof.
7. The composition of Claim 6 wherein the liquid carrier comprises from about 40% to about 90% by weight of water and from about 1% to about 15% by weight of the C₁-C₆ alkanols.
8. The composition of Claim 7 wherein the C₁-C₆ alkanols are selected from the group consisting of ethanol, n-propanol, isopropanol, n-butanol, amyl alcohol, and mixtures thereof.
9. The composition of Claim 2 wherein the propellant is selected from the group consisting of propane, butane, isobutane, nitrogen, carbon dioxide, nitrous oxide, atmospheric gas, 1,2-difluoroethane, dimethylether, and mixtures thereof.

10. The composition of Claim 2 wherein the composition further comprises a combination of the polyalkylene glycol and a polysaccharide styling polymer.
11. The composition of Claim 10 wherein the polysaccharide styling polymer is selected from the group consisting of glucosamine polysaccharide derivatives, cationic polysaccharides, anionic polysaccharides, and mixtures thereof.
12. The composition of Claim 2 wherein the composition further comprises a gelling agent.
13. The composition of Claim 12 wherein the gelling agent is a water-soluble polymer selected from the group consisting of crosslinked ethylene/maleic anhydride copolymers, crosslinked carboxylic acid polymers, hydrophobically modified nonionic cellulose polymers, and mixtures thereof.
14. A hair mousse composition comprising:
 - (a) from about 5% to about 25% by weight of a polyalkylene glycol that is substantially free of polyalkylene glyceryl ethers and that has a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms;
 - (b) from about 10% to about 90% by weight of water; and
 - (c) from about 5% to about 40% by weight of a propellant.
15. The composition of Claim 14 wherein the polyalkylene glycol has a number average molecular weight of from about 400 to about 1500.
16. The composition of Claim 15 wherein the polyalkylene glycol is selected from the group consisting of ethoxy polyethylene/polypropylene glycol copolymers, methoxy polyethylene/polypropylene glycol copolymers, propoxy polyethylene/polypropylene glycol copolymers, butoxy polyethylene/polypropylene glycol copolymers, pentoxy polyethylene/polypropylene glycol copolymers, triglycerin, hexaglycerin, PPG-4, PPG-6, PEG-5, PEG-6, PEG-8, PEG-12, PEG-14, PEG-18, PEG-20, PEG-32, and mixtures thereof.
17. The composition of Claim 16 wherein the polyalkylene glycol is selected from the group consisting of PPG-4, PEG-8, PEG-12, PEG-20, and mixtures thereof.

18. The composition of Claim 14 wherein the propellant is selected from the group consisting of propane, butane, isobutane, nitrogen, carbon dioxide, nitrous oxide, atmospheric gas, 1,2-difluoroethane, dimethylether, and mixtures thereof.
19. The composition of Claim 14 wherein the composition further comprises an organic solvent selected from the group consisting of C₁-C₆ alkanols, carbitol, acetone, and mixtures thereof.
20. The composition of Claim 19 wherein the composition comprises from about 40% to about 90% by weight of water and from about 1% to about 15% by weight of the organic solvent.
21. The composition of Claim 20 wherein the organic solvent is a C₁-C₆ alkanol selected from the group consisting of ethanol, n-propanol, isopropanol, n-butanol, amyl alcohol, and mixtures thereof.
22. The composition of Claim 14 wherein the composition further comprises a combination of the polyalkylene glycol and a polysaccharide styling polymer.
23. The composition of Claim 22 wherein the polysaccharide styling polymer is selected from the group consisting of glucosamine polysaccharide derivatives, cationic polysaccharides, anionic polysaccharides, and mixtures thereof.
24. The composition of Claim 14 wherein the composition further comprises a gelling agent.
25. The composition of Claim 24 wherein the gelling agent is a water-soluble polymer selected from the group consisting of crosslinked ethylene/maleic anhydride copolymers, crosslinked carboxylic acid polymers, hydrophobically modified nonionic cellulose polymers, and mixtures thereof.

26. A method of making an improved hair mousse composition wherein the method comprises the steps of:

(a) preparing a reformable weld formulation comprising:

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(i) from about 9% to about 25% by weight of a polyalkylene glycol having a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms;

(ii) from about 40% to about 90% by weight of water; and

(iii) from about 1% to about 15% by weight of an organic solvent; and

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(b) packaging the reformable weld formulation in an aerosol container, wherein the aerosol container comprises a propellant, and wherein the reformable weld formulation is substantially free of polyalkylene glyceryl ethers.

27. The method of Claim 26 wherein the reformable weld formulation comprises from about 10% to about 20% by weight of the polyalkylene glycol.

28. The method of Claim 27 wherein the polyalkylene glycol has a number average molecular weight of from about 400 to about 1500.

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29. The method of Claim 28 wherein the polyalkylene glycol is selected from the group consisting of ethoxy polyethylene/polypropylene glycol copolymers, methoxy polyethylene/polypropylene glycol copolymers, propoxy polyethylene/polypropylene glycol copolymers, butoxy polyethylene/polypropylene glycol copolymers, pentoxy polyethylene/polypropylene glycol copolymers, triglycerin, hexaglycerin, PPG-4, PPG-6, PEG-5, PEG-6, PEG-8, PEG-12, PEG-14, PEG-18, PEG-20, PEG-32, and mixtures thereof.

30. The method of Claim 29 wherein the polyalkylene glycol is selected from the group consisting of PPG-4, PEG-8, PEG-12, PEG-20, and mixtures thereof.

31. The method of Claim 26 wherein the organic solvent is selected from the group consisting of C₁-C₆ alkanols, carbitol, acetone, and mixtures thereof.

32. The method of Claim 31 wherein the organic solvent is a C₁-C₆ alkanol selected from the group consisting of ethanol, n-propanol, isopropanol, n-butanol, amyl alcohol, and mixtures thereof.

33. The method of Claim 26 wherein the propellant is selected from the group consisting of propane, butane, isobutane, nitrogen, carbon dioxide, nitrous oxide, atmospheric gas, 1,2-difluoroethane, dimethylether, and mixtures thereof.

34. A hair styling composition comprising:

- (a) from about 65% to about 99% by weight of a polyalkylene glycol that is substantially free of polyalkylene glyceryl ethers and that has a number average molecular weight of from about 190 to about 1500 and from about 5 to about 35 repeating alkylene oxide radicals wherein each of the repeating alkylene oxide radicals has from 2 to 6 carbon atoms; and
- (b) from about 1% to about 30% by weight of a liquid carrier.

35. The composition of Claim 34 wherein the polyalkylene glycol has a number average molecular weight of from about 400 to about 1500.

36. The composition of Claim 35 wherein the polyalkylene glycol is selected from the group consisting of ethoxy polyethylene/polypropylene glycol copolymers, methoxy polyethylene/polypropylene glycol copolymers, propoxy polyethylene/polypropylene glycol copolymers, butoxy polyethylene/polypropylene glycol copolymers, pentoxy polyethylene/polypropylene glycol copolymers, triglycerin, hexaglycerin, PPG-4, PPG-6, PEG-5, PEG-6, PEG-8, PEG-12, PEG-14, PEG-18, PEG-20, PEG-32, and mixtures thereof.

37. A method for styling dry hair, which method comprises applying an effective amount of the composition of Claim 1 to the hair.

38. A method for styling dry hair, which method comprises applying an effective amount of the composition of Claim 14 to the hair.

39. A method for styling dry hair, which method comprises applying an effective amount of the composition of Claim 26 to the hair.

40. A method for styling dry hair, which method comprises applying an effective amount of the composition of Claim 34 to the hair.

[illegible][illegible]

DECLARATION COMBINED WITH POWER OF ATTORNEY

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Attorney Docket No. 7547

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled AEROSOL HAIR STYLING COMPOSITIONS CONTAINING SELECT POLYALKYLENE GLYCOL STYLING AGENTS the specification of which

☒ is attached hereto.
☐ was filed on _____ as United States Application No. _____ or
PCT International Application Number _____
and was amended on _____ (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37 Code of Federal Regulations §1.56.

I hereby claim foreign priority benefits under Title 35 United States Code §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate, or §365(a) of any PCT International application which designated at least one country other than the United States of America, listed below and have also identified below any foreign application for patent or inventor's certificate, or of any PCT international application having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s)			Priority Claimed	
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No
_____ (Number)	_____ (Country)	_____ (Day/Month/Year Filed)	<input type="checkbox"/> Yes	<input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code §119(e) of any United States provisional application(s) listed below.

Application Serial No.	Filing Date	Application Serial No.	Filing Date
------------------------	-------------	------------------------	-------------

I hereby claim the benefit under Title 35, United States Code §120 of any United States application(s), or §365(c) of any PCT International application designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code §112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

U.S. Parent Application Number	PCT Parent Number	Parent Filing Date (MM/DD/YYYY)	Parent Patent Number (If applicable)

As named inventor, I hereby appoint the following registered practitioner(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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